



## UNITED STATES ENVIRONMENTAL PROTECTION AGENCY

WASHINGTON, D.C. 20460

OFFICE OF  
PREVENTION, PESTICIDES  
AND TOXIC SUBSTANCES

Diane Allemang  
Registrant Representative  
Jellenek, Schwartz & Connolly, Inc.  
1525 Wilson Boulevard  
Suite 600  
Arlington, VA 22209

Dear Ms. Allenmang:

This letter is to transmit Office of Pesticide Programs' (OPP) preliminary human health and ecological risk assessments for the organophosphate (OP) ethyl parathion, for which the company you represent holds a registration for technical material. We are providing you with a 30-day period to identify and comment on errors only. Your comments must be received within 30 days of receipt of this letter. The Agency will review and evaluate your comments on errors upon receipt. On or about January 4, 1999, the preliminary risk assessments, your comments and the Agency's review and discussion of your comments will be placed into OPP's Public Docket. The docket will be opened for public comment for 60 days from the date of publication of a Notice of Data Availability in the *Federal Register*. Additionally, the preliminary risk assessments will be placed on the internet at the same time. This process is part of the Agency's efforts to involve the public in the implementation of the Food Quality Protection Act of 1996 and serves as an interim measure to improve the transparency of the reregistration and tolerance reassessment processes.

During this 30-day comment period, the Agency asks for comments on errors, confidential business information (CBI), and planned data. The Agency will respond only to errors which do not pertain to matters of policy, interpretation, or applicability of data. Errors include, but are not limited to, mathematical, computational, typographic, or other similar errors. In the process of reviewing the Agency's preliminary risk assessments, we ask that you inform us in writing of any claims of CBI contained in these assessments. If we do not receive notification in writing of any such claims within 30 days, the Agency will assume that the documents are free of CBI. Also, we request that you inform the Agency of any pertinent, on-going or planned studies, or other sources of information on ethyl parathion, and your timetable for completing and submitting such data and information to the Agency. This will enable the Agency to plan better for refining the risk assessments and completing the reregistration and tolerance reassessment.

If during this thirty day period, you submit comments other than on errors, you should clearly indicate that they are submitted in advance for the 60-day public comment period. You

should provide a brief summary of the comments and refer to an attachment which provides a more in-depth discussion.

Please mail your response, any claims of CBI and other information about ethyl parathion to Dennis Deziel. In addition, provide Mr. Deziel your response and any supporting material in both hard copy and electronic form. If you have any questions, please contact him at (703) 308-8173.

Sincerely,

Arnold E. Layne, Chief  
Reregistration Branch 1  
Special Review and  
Reregistration Division